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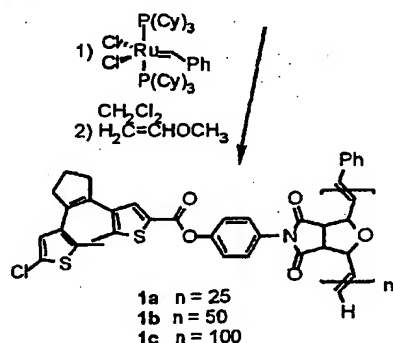
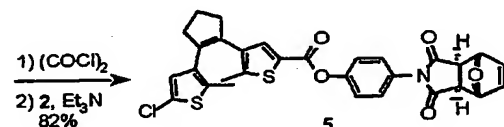
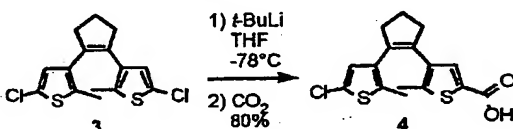
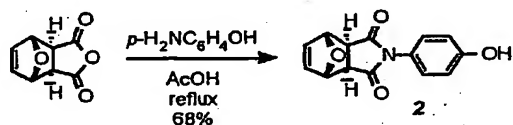
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(54) Title: NOVEL PHOTOCHROMIC POLYMERS AND METHODS OF SYNTHESIZING SAME



Preparation of Homopolymers 1a-1c

(57) Abstract: Ring-opening metathesis polymerization (ROMP) of a photochromic 1,2-bis-(3-thienyl)-cyclopentene monomer generated a series of novel polymers. All polymers exhibit reversible light-activated interconversion between their colorless-open and their colored-closed forms.

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## NOVEL PHOTOCHROMIC POLYMERS AND METHODS OF SYNTHESIZING SAME

### FIELD OF THE INVENTION

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The present invention relates to novel photochromic polymers, methods of preparing the photochromic polymers and all uses of the novel photochromes for example in optical materials and photonic devices.

### 10 BACKGROUND OF THE INVENTION

15 The photochromism of 1,2-bis-(3-thienyl)-cyclopentene derivatives involves the reversible photoinduced cyclization between the colorless-open and the colored-closed forms of the chromophore<sup>1</sup>. This phenomenon is interesting for its potential role in optical materials and photonic devices such as variable-transmission filters, optical information storage systems and photo-regulated molecular switches<sup>2</sup>. The need for practical handling of definite forms of photochromic materials such as films, sheets, fibers or beads, dictate the use of polymeric rather than monomeric photochromes<sup>3,4</sup>. Homopolymers (i.e. polymers derived from one species of monomer) are more desirable than copolymers as they will have an increased density of the photochromic unit within the material. This translates into a greater amount of information expressed or stored per unit volume or surface.

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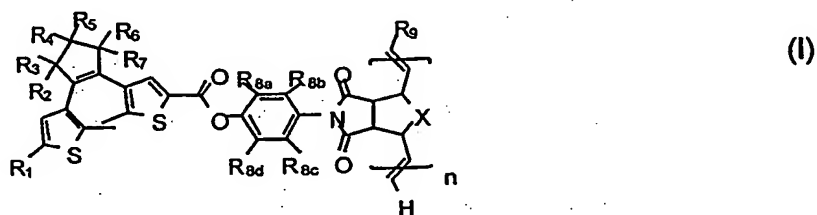
Existing polymerization processes rely on harsh reaction conditions which limits the structure of photochromic products. The present invention employs very mild reaction conditions which allows compatibility and flexibility, including the capacity to synthesize products having a variety of pendant functional groups. The chemistry can be controlled and therefore macroscopic properties may be tailored.

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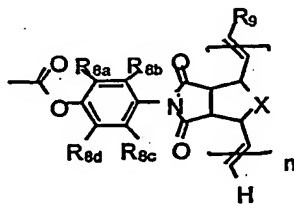
In view of the foregoing, there is a need in the art to develop novel photochromic homopolymers and methods of synthesizing same.

SUMMARY OF THE INVENTION

The present invention relates to novel homopolymers of the  
Formula I:



10 wherein  $n$  is any integer;  $X$  is  $\text{CH}_2$ ,  $\text{CHR}_{10}$ ,  $\text{O}$ ,  $\text{S}$ ,  $\text{NH}$ ,  $\text{N-alkyl}$ ,  $\text{N-aryl}$  or  $\text{CH}_2\text{-CH}_2$ ;  $R_1$  is selected from the group consisting of  $\text{H}$ , a halogen, a carboxylic acid, an ester, an aldehyde, an alkene, a phenyl, an aryl and structure A where structure A is:



20  $R_2$ ,  $R_3$ ,  $R_4$ ,  $R_5$ ,  $R_6$  and  $R_7$  are each selected from the group consisting of  $\text{H}$  and a halogen;  $R_{8a}$ ,  $R_{8b}$ ,  $R_{8c}$  and  $R_{8d}$  are each selected from the group consisting of  $\text{H}$ , an alkyl and a halogen;  $R_9$  is a phenyl or an aryl; and  $R_{10}$  is an alkyl, an aryl or a phenyl. Preferably,  $n$  is from about 10 to about 100,  $R_1$  is chlorine, phenyl or structure A,  $R_2 - R_7$  is  $\text{H}$  or  $\text{F}$ ,  $R_{8a} - R_{8d}$  is  $\text{H}$ ,  $R_9$  is phenyl  
25 and  $X$  is  $\text{O}$  or  $\text{CH}_2$ .

The novel photochromic homopolymers of the invention are prepared using ring-opening metathesis polymerization (ROMP) of a photochromic 1,2-*bis*-(3-thienyl)-cyclopentene monomer. All novel polymers  
30 of the invention exhibit reversible light-activated interconversion between their colorless-open and their colored-closed forms.

**BRIEF DESCRIPTION OF THE DRAWINGS**

Figure 1 is a schematic drawing showing the preparation of photochromic homopolymer 1 of the invention.

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Figures 2(a) and 2(b) are graphs showing changes in the UV-Vis absorption spectra of THF solutions of (a) monomer 5 ( $3 \times 10^{-5}$  M) and (b) polymer 1a ( $3 \times 10^{-5}$  M with respect to the 1,2-bis-(3-thienyl)-cyclopentene photochrome) upon irradiation with 254 nm light. Irradiation periods are 2, 4,

10 6, 8, 10, 15, 20, 25, 30 and 40 seconds.

Figure 3 is a schematic drawing showing the preparation of photochromic homopolymer 9 of the invention.

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Figure 4 is a schematic drawing showing the preparation of photochromic homopolymer 12 of the invention.

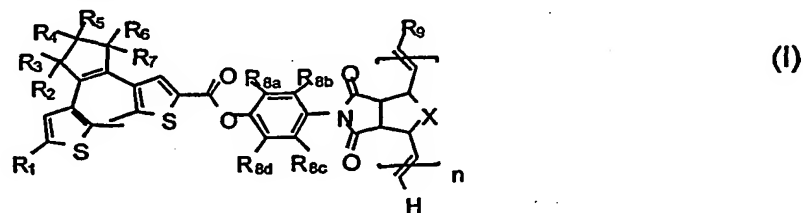
Figure 5 is a schematic drawing showing the preparation of photochromic homopolymer 17 of the invention.

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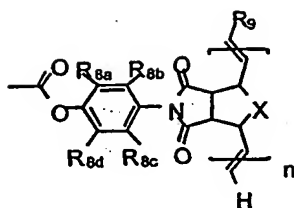
**DETAILED DESCRIPTION OF THE INVENTION**

As hereinbefore mentioned, the present invention relates to novel photochromic homopolymers of the Formula I:

25



30 wherein n is any integer; X is CH<sub>2</sub>, CHR<sub>10</sub>, O, S, NH, N-alkyl, N-aryl or CH<sub>2</sub>-CH<sub>2</sub>; R<sub>1</sub> is selected from the group consisting of H, a halogen, a carboxylic acid, an ester, an aldehyde, an alkene, a phenyl, an aryl and structure A where structure A is:

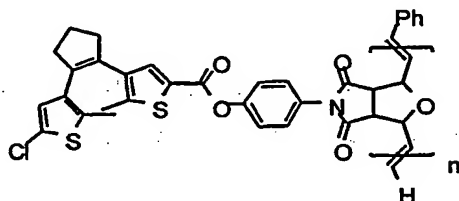


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$R_2$ ,  $R_3$ ,  $R_4$ ,  $R_5$ ,  $R_6$  and  $R_7$  are each selected from the group consisting of H and a halogen;  $R_{8a}$ ,  $R_{8b}$ ,  $R_{8c}$  and  $R_{8d}$  are each selected from the group consisting of H, an alkyl and a halogen;  $R_9$  is a phenyl or an aryl; and  $R_{10}$  is an

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Preferably, the homopolymer is of the Formula II



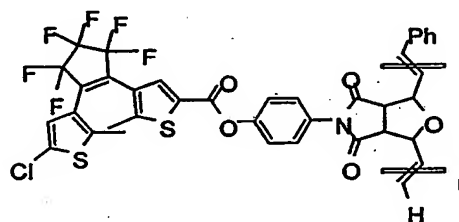
(II)

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wherein  $n$  is from about 10 to about 100.

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Other preferred homopolymers of the present invention include the fluorinated homopolymer of Formula III,



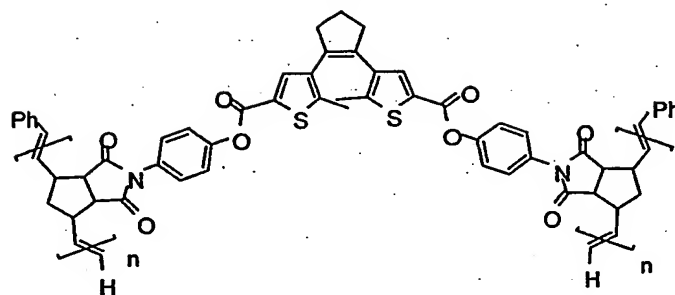
(III)

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the cross-linked homopolymer of Formula IV,

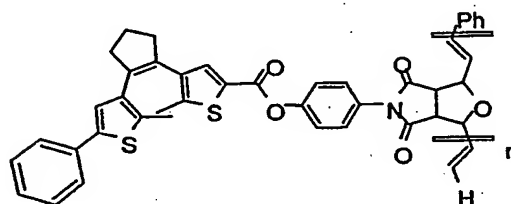
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(IV)

and the phenyl homopolymer of Formula V



(V)

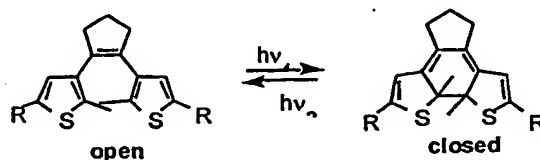
wherein  $n$  is from about 10 to about 100 for each of Formulas III, IV and V.

The homopolymers of the invention are prepared by ring-opening metathesis polymerization (ROMP) of a photochromic 1,2-bis-(3-thienyl)-cyclopentene monomer. The preparation of homopolymers of the Formulas II, III, IV and V are described in detail in Examples 1, 2, 3 and 4 below and are shown in Figures 1, 3, 4 and 5, respectively. The ROMP process is also described in U.S. Patent No. 5,849,851 which is incorporated herein by reference.

Photochromes based on 1,2-bis-(3-thienyl)-cyclopentene derivatives offer many advantages. For example 1,2-bis-(3-thienyl)-cyclopentene derivatives possess optimal photochromic properties, including thermal irreversibility and fatigue resistance. In addition, the wavelength of light expressed in the colored forms can be readily tuned by tailoring the electronic distribution in the conjugated pathway created upon cyclization. This is most conveniently accomplished by modifying pendant functional groups (R) located on the heterocycles as shown below in the reaction

scheme that illustrates the photoinduced interconversion between the colorless-open and colored-closed form of the 1,2-bis-(3-thienyl)-cyclopentene photochrome.

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The inventors determined that in order to take full advantage of all possible pendant groups so that a wider range of colors can be expressed by the polymers, a mild and tolerant reaction polymerization protocol must be found. The inventors further determined that ring-opening metathesis polymerization (ROMP) techniques<sup>5</sup> provide the desired conditions and thus do not limit the pool of pendant R groups that can be used. As a result, the R group can be any functional group including, but not limited to, a halogen, a carboxylic acid, an ester, an aldehyde, an alkene or structure A.

Advantageously, the ROMP process produces homopolymers rather than copolymers. This provides macroscopic materials that have an increased density of photochromic units, a feature that is very advantageous for practical applications. Another advantage of using the ROMP technology is that by varying the catalyst-to-substrate stoichiometry, ROMP allows the polymer chain length to be readily and precisely tailored.

The present application represents the first example of photochromic homopolymers synthesized using ROMP technology<sup>6</sup>. The result is an unprecedented series of well-ordered homogeneous homopolymers that reversibly undergo changes in color when irradiated with light of appropriate wavelength. Color changes are, for example, colorless-to-red or colorless-to-purple. These colors can be easily tuned by simple tailoring of the photochrome's pendant groups.

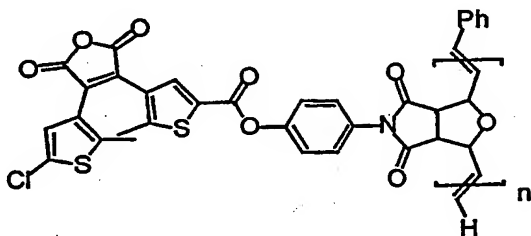
The present invention includes all uses of the novel homopolymers of the inventions. These uses include, but are not limited to,



- (1) ophthalmic lenses-eyeglasses that change color depending on the ambient light;
- (2) actinometry, and molecular sensors;
- (3) novelty items such as photochromic inks, paints and fibers;
- 5 (4) variable transmission filters — those that on command, regulate the amount and type of light that can be transmitted;
- (5) high-density optical information storage systems (this invention is particularly well-suited to this application as it provides more information storage sites per unit area);
- 10 (6) photo-regulated molecular switches that can be incorporated into molecular-scale machinery;
- (7) optoelectronic systems; and
- (8) reversible holographic systems.

15 As will be appreciated by a person skilled in the art, co-polymers or other polymers comprising the photochromic units of the present invention may also readily be produced utilizing ROMP technology. Such other polymers are within the scope of the present invention.

20 In other embodiments of the invention, the polymerization protocol described herein, including ROMP techniques and the use of Grubbs' catalyst, may be used in connection with other photochromic monomers, including an anhydride variant to derive the anhydride polymer shown below:



25

## EXAMPLES

The following examples will further illustrate the invention in greater detail although it will be appreciated that the invention is not limited to the specific examples.

### Experimental Methods

Tetrahydrofuran (THF) (Caledon) was distilled over sodium/benzophenone ketyl radical under argon.  $\text{CH}_2\text{Cl}_2$  (Caledon) was distilled over calcium hydride under argon. All other solvents were used as received. Solvents for NMR analysis (Cambridge Isotope Laboratories) were used as received. Bis(tricyclohexylphosphine)benzylidene ruthenium(IV)dichloride (Grubbs' catalyst)<sup>7</sup> was purchased from Strem and was stored and weighed in a glove box under a nitrogen atmosphere. All other reagents and starting materials were purchased from Aldrich. Compounds 7-oxa-bicyclo[2.2.1]hept-5-ene-2,3-dicarboxylic anhydride<sup>8</sup>, **2a**<sup>9</sup>, **3**<sup>10</sup>, **6**<sup>11</sup>, and **10**<sup>12</sup> were prepared as described in the literature and as would be readily apparent to a person skilled in the art. Compound **13** was prepared in a fashion analogous to the synthesis of compound **4** except that carbon dioxide was replaced with iodine as would be readily apparent to a person skilled in the art.

All polymerization reactions were carried out on a Schlenk line. Gel permeation chromatography (GPC) analyses (calibrated by polystyrene) were performed on THF solutions of the polymers using a Waters 515 HPLC pump and 2410 Refractive Index Detector at a flow rate of 1.0 mL/minute through a 7.8 x 300 mm column at 30°C. <sup>1</sup>H NMR characterizations were performed on a Varian Inova-300 instrument, working at 299.96 MHz. Chemical shifts ( $\delta$ ) are reported in parts per million relative to tetramethylsilane using the residual solvent peak as a reference standard. Coupling constants ( $J$ ) are reported in Hertz. FT-IR measurements were

performed using a Nicolet Magna-IR 750. UV-Vis measurements were performed using a Pharmacia Biotech Ultraspec 3000 spectrophotometer.

### Preparation of Homopolymers

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#### Example 1 – Preparation of Homopolymer 1 (Figure 1)

##### **Step 1 – Synthesis of Olefin fragment 2**

A mixture of 7-oxa-bicyclo[2.2.1]hept-5-ene-2,3-dicarboxylic anhydride (2.0 g, 12 mmol) and *p*-aminophenol (1.31 g, 12 mmol) were heated at reflux for 10 minutes in glacial acetic acid (3 mL), after which time a precipitate formed. The reaction mixture was cooled to room temperature and the product was collected by filtration, washed with water and dried *in vacuo* to afford 2.1 g (68%) of the olefin fragment *exo*-N-(*p*-hydroxyphenyl)-3,6-epoxy-4-cyclohexene-1,2-dicarboximide **2** as a white solid: <sup>1</sup>H NMR (DMSO-*d*<sub>6</sub>) δ 9.71 (s, 1H), 6.95 (d, *J* = 8.75 Hz, 2H), 6.83 (d, *J* = 8.75 Hz, 2H), 6.58 (s, 2H), 5.21 (s, 2H), 3.02 (s, 2H); <sup>13</sup>C NMR (DMSO-*d*<sub>6</sub>) δ 175.95, 157.27, 136.53, 127.98, 123.19, 115.38, 80.66, 47.20; EIMS (*m/z*): 257 (*M*<sup>+</sup>), 189 (*M*<sup>+</sup> - C<sub>4</sub>H<sub>4</sub>O); FTIR  $\bar{\nu}$  3334 (s, broad), 3143, 3102, 3076, 3049, 3029, 2973, 1697, 1612, 1594.

##### **Step 2 – Synthesis of carboxylic acid 4**

A solution of dichloride **3** (385 mg, 1.17 mmol) in dry THF (50 mL) at -78°C under argon was treated with *tert*-butyllithium (0.7 mL of 1.7 M solution in hexane, 1.17 mmol). After stirring for 15 minutes, excess dry CO<sub>2</sub> was bubbled through the solution. The reaction mixture was warmed to room temperature, quenched with dilute HCl, extracted with Et<sub>2</sub>O (3 x 50 mL), dried over Na<sub>2</sub>SO<sub>4</sub> and evaporated *in vacuo*. Purification by column chromatography through silica (5% CH<sub>3</sub>OH-CH<sub>2</sub>Cl<sub>2</sub>) afforded 318 mg (80%) of the product **4** as a pale yellow solid: <sup>1</sup>H NMR (CDCl<sub>3</sub>) δ 7.55 (s, 1H), 6.52 (s, 1H), 2.72 (m, 4H), 1.99 (m, 5H), 1.80 (s, 3H).

The yield of the monoanion in the lithiation step is greater than would be expected from a statistically governed product distribution. This may be attributed to the unfavorable charge build-up that would exist if both chlorine atoms underwent metal-halogen exchange reactions. This charge build-up is still significant despite the fact that the two thiophene rings are cross-conjugated instead of directly conjugated, suggesting the charge-charge repulsion between two thiophene heterocycles would be felt through space.

#### 10 Step 3 – Synthesis of monomer 5

A vigorously stirred solution of carboxylic acid 4 (136 mg, 0.4 mmol) and 5 drops of DMF in  $\text{CH}_2\text{Cl}_2$  (4 mL) at  $0^\circ\text{C}$  was treated with a solution of oxalyl chloride (254 mg, 2.0 mmol) in  $\text{CH}_2\text{Cl}_2$  (6 mL) dropwise over 10 minutes. After stirring at room temperature under argon for 2 hours, the reaction mixture was concentrated to dryness *in vacuo*. The residue was dissolved in  $\text{CH}_2\text{Cl}_2$  (10 mL) and added dropwise over 10 minutes to a solution of olefin 2 (154 mg, 0.6 mmol) and triethylamine (0.5 mL) in acetone cooled to  $0^\circ\text{C}$ . The mixture was stirred overnight under argon and the solvent was removed *in vacuo*. Purification by column chromatography through silica (2%  $\text{CH}_3\text{OH}-\text{CHCl}_3$ ) afforded 188 mg (82%) of the product as a pale yellow solid:  $^1\text{H}$  NMR ( $\text{CDCl}_3$ )  $\delta$  7.67 (s, 1H), 7.31 (m, 4H), 6.56 (s, 3H), 5.38 (m, 2H), 3.00 (s, 2H), 2.76 (m, 4H), 2.05 (m, 5H), 1.87 (s, 3H);  $^{13}\text{C}$  NMR ( $\text{CDCl}_3$ )  $\delta$  175.22, 159.95, 150.56, 144.60, 137.19, 136.78, 136.02, 135.33, 134.75, 134.14, 133.36, 129.16, 128.03, 127.62, 126.66, 125.59, 122.38, 81.51, 47.59, 38.61, 38.52, 22.89, 14.98, 14.30; ESMS(+ive): 600.0 ( $\text{M}+\text{Na}^+$ ), 532 ( $\text{M}-\text{Cl}$ ); IR (KBr-cast): 3050 (w), 2951 (w), 2843 (w), 1713 (s), 1202 (s).

#### Step 4 – Polymerization of bicyclic olefin 5 to yield polymers 1(a)-(c)

ROMP reactions of monomer 5 were performed under rigorously inert conditions in a Schlenk tube. In a typical procedure a solution of monomer 5 in dry deoxygenated  $\text{CH}_2\text{Cl}_2$  was cannulated into a  $\text{CH}_2\text{Cl}_2$  solution of bis(tricyclohexylphosphine)benzylidene ruthenium(IV)dichloride (Grubbs' catalyst) (0.04 equivalents for 1a, 0.02 equivalents for 1b, and 0.01

equivalents for **1c**). The final monomer concentrations were 0.1 M. After stirring at room temperature for 14 hours, excess ethylvinyl ether was added and the solutions were stirred exposed to the atmosphere for 30 minutes. The polymers were precipitated in pure form by pouring the reaction solutions into cold Et<sub>2</sub>O and collecting the precipitates by vacuum filtration (yields: 75% for **1a**, 78% for **1b**, and 75% for **1c**). Homopolymers with varying molecular weights were synthesized in a systematic fashion by changing the molar amount (1-4 mol%) of Grubbs' catalyst used to initiate the reaction. In all cases, the ROMP reactions were reproducible, affording polymers **1a-c** as off-white powders in good yields (~75%).

The success of the polymerization reactions was assessed by using <sup>1</sup>H NMR spectroscopy. In the <sup>1</sup>H NMR spectrum of monomer **5**, the peak corresponding to the methine proton of the bicyclic olefin overlaps with the peak corresponding to the hydrogen on the chlorothiophene heterocycle at 6.56 ppm resulting in the area under the peak integrating to 3 protons. After the polymerization reactions, the peak at 6.56 ppm integrated to only 1 proton with respect to the other thiophene proton at 7.67 ppm. A new peak for the olefin protons appeared at 6.10 ppm. A typical <sup>1</sup>H NMR (CDCl<sub>3</sub>) is as follows: δ 7.6 (br s), 7.3 (br s), 6.6 (br s), 6.1 (br s), 5.8 (m), 5.2 (m), 4.6 (m), 3.4 (br s), 2.7 (br s), 2.0 (m), 1.8 (br s). The GPC characterization of the polymeric products is set forth in Table 1.

The polymeric products were all readily soluble in common organic solvents such as chloroform, dichloromethane, tetrahydrofuran and benzene. They are air-stable solids of reasonable number-average molecular weights (M<sub>n</sub>) and relatively narrow polydispersities (M<sub>w</sub>/M<sub>n</sub>) as determined by gel permeation chromatographic analyses calibrated by polystyrene (Table 1).

Table 1 also summarizes the UV-Vis absorption properties of THF solutions of the novel polymers along with those for monomers **5**, **8**, **11** and **16**. Polymers **1a - c**, for example, show typical absorbances for the colorless-open forms of the 1,2-bis-(3-thienyl)-cyclopentene photochrome at

248 nm. Photo-induced isomerization studies were carried out by irradiating the THF solutions of 1a-c at 254 nm with a short-wavelength hand held lamp. Spectral changes were monitored in the UV-Vis region (Figure 2). In all cases, irradiation produced an immediate decrease in the absorbances corresponding to the open form of the dithienylethene photochrome at 248 nm. The decrease in these absorbances was accompanied by the appearance of broad absorbances for the pink closed form of the photochrome centered at 512 nm. A typical irradiation time appropriate to effect quantitative ring closure is 40 seconds for solutions at  $3 \times 10^{-5}$  M concentrations. The similarity of the absorption spectrum (values of  $\lambda_{\max}$  and  $\epsilon$ ) for monomer 5 and those of the polymers in both open and closed forms illustrates that the intimacy of the photochromes covalently linked to the polymer backbone affects neither the ground-state nor the excited-state properties of the photochrome. As shown in Table 1, similar results were found in respect of the other homopolymers of this invention.

The polymers can be easily decolorized by subsequent irradiation for 1 minute with light of wavelength greater than 434 nm using a high power-lamp with an appropriate cutoff filter. The photoisomerization process is thermally irreversible, and solutions of the closed forms kept in the dark showed no changes in their absorption spectra.

The polymers were very robust, and their thin films were easily cast by layering toluene solutions of 1a-c onto water. After slow evaporation of the toluene, the resulting transparent films were transferred to slides.

Transmission electron microscopy of the films reveals a smooth surface possessing no topological features. These films immediately turned from colorless to pink when exposed for brief periods (60 seconds) at 254 nm. As with the polymers in solution, the films could be decolorized with above 434 nm light.

**Example 2 – Preparation of Flourinated Homopolymer 9 (Figure 3)****Step 1 – Synthesis of carboxylic acid 7**

A solution of dichloride 6 (500 mg, 1.14 mmol) in dry THF (50 mL) at -78°C under argon was treated with *tert*-butyllithium (0.68 mL of 1.7 M solution in hexane, 1.14 mmol). After stirring for 15 minutes, excess dry CO<sub>2</sub> was bubbled through the solution. The reaction mixture was warmed to room temperature, quenched with dilute HCl, extracted with Et<sub>2</sub>O (3 x 50 mL), dried over Na<sub>2</sub>SO<sub>4</sub> and evaporated *in vacuo*. Purification by column chromatography through silica (5% CH<sub>3</sub>OH-CH<sub>2</sub>Cl<sub>2</sub>) afforded 260 mg (51%) of the product as a white solid: <sup>1</sup>H NMR (CDCl<sub>3</sub>) δ 7.83 (s, 1H), 6.85 (s, 1H), 2.02 (s, 3H), 1.83 (s, 3H).

**Step 2 – Synthesis of monomer 8**

A vigorously stirred solution of carboxylic acid 7 (254 mg, 0.57 mmol) and 5 drops of DMF in CH<sub>2</sub>Cl<sub>2</sub> (8 mL) at 0°C was treated with a solution of oxalyl chloride (361 mg, 2.8 mmol) in CH<sub>2</sub>Cl<sub>2</sub> (4 mL) dropwise over 10 minutes. After stirring at room temperature under argon for 2 hours, the reaction mixture was concentrated to dryness *in vacuo*. The residue was dissolved in CH<sub>2</sub>Cl<sub>2</sub> (8 mL) and added dropwise over 10 minutes to a solution of olefin 2 (221 mg, 0.86 mmol) and triethylamine (0.5 mL) in acetone cooled to 0°C. The mixture was stirred overnight under argon and the solvent was removed *in vacuo*. Purification by column chromatography through silica (CHCl<sub>3</sub>) afforded 75% yield of the product as a pale yellow solid: <sup>1</sup>H NMR (CDCl<sub>3</sub>) δ 7.91 (s, 1H), 7.33 (m, 4H), 6.88 (s, 1H), 6.56 (m, 2H), 5.39 (m, 2H), 3.01 (s, 2H), 2.04 (s, 3H), 1.88 (s, 3H); <sup>13</sup>C NMR (CDCl<sub>3</sub>) δ 175.20, 159.12, 150.19, 140.64, 136.79, 134.56, 130.66, 129.50, 128.40, 127.74, 126.02, 125.48, 123.91, 122.25, 81.52, 47.59, 15.08, 14.52; ESMS(+ive): 703.1 (M+NH<sub>4</sub><sup>+</sup>, 100%); IR (KBr-cast): 3015 (w), 1714 (s).

**Step 3 – Polymerization of bicyclic olefin 8 to yield polymer 9**

The procedure is identical to that for the polymerization of olefin 5 except that the polymer was precipitated by adding MeOH instead of Et<sub>2</sub>O.

Thus **8** (37 mg, 0.056 mmol) was polymerized with Grubbs' catalyst (1.8 mg, 0.04 equivalents) to yield polymer **9** in 53% yield:  $^1\text{H}$  NMR ( $\text{CDCl}_3$ )  $\delta$  7.9 (br s), 7.3 (br s), 6.9 (br s), 6.1 (br s), 5.8 (m), 5.2 (m), 4.6 (m), 3.4 (br s), 2.0 (m), 1.8 (br s). GPC:  $M_n=7437$ ,  $M_w=10953$ , Polydispersity Index=1.47. Polymer **9** exhibited a color change from colorless to purple upon irradiation with 254-nm light.

### Example 3 – Preparation of Cross-linked homopolymer **12** (Figure 4)

#### 10 Step 1 – Synthesis of diolefin monomer **11**

A vigorously stirred solution of dicarboxylic acid **10** (375 mg, 1.08 mmol) and DMF (0.1 ml) in  $\text{CH}_2\text{Cl}_2$  (20 mL) at  $0^\circ\text{C}$  was treated with a solution of oxalyl chloride (2.02 g, 16 mmol) in  $\text{CH}_2\text{Cl}_2$  (10 mL) dropwise over 10 minutes. After stirring at room temperature under argon for 2 hours, the reaction mixture was concentrated to dryness *in vacuo*. The residue was dissolved in  $\text{CH}_2\text{Cl}_2$  (20 mL) and added dropwise over 10 minutes to a solution of olefin **2a** (833 mg, 3.24 mmol) and triethylamine (2 mL) in acetone cooled to  $0^\circ\text{C}$ . The mixture was stirred overnight under argon and the solvent was removed *in vacuo*. Purification by column chromatography through silica (1.5% MeOH- $\text{CHCl}_3$ ) afforded 400 mg (45% yield) of the product as a pale pink solid:  $^1\text{H}$  NMR ( $\text{CDCl}_3$ )  $\delta$  7.66 (s, 2H), 7.22 (m, 8H), 6.23 (m, 4H), 3.48 (m, 4H), 3.41 (m, 4H), 2.81 (m, 4H), 2.08 (m 2H), 1.99 (s, 6H), 1.76, (m, 2H), 1.58 (d,  $J=8.7$  Hz, 2H);  $^{13}\text{C}$  NMR ( $\text{CDCl}_3$ )  $\delta$  176.64, 159.90, 150.31, 144.43, 137.01, 135.89, 134.95, 134.68, 129.35, 128.30, 127.67, 122.27, 52.30, 45.83, 45.58, 38.75, 22.88, 15.07; ESMS(+ive): 845.2 ( $M+\text{Na}^+$ , 100%); IR (KBr-cast): 2949 (w), 1709 (s).

#### Step 2 – Polymerization of bis-bicyclic olefin **11** to yield homopolymer **12**

The procedure is identical to that for the polymerization of olefin **5**. Thus **11** (81mg, 0.098 mmol) was polymerized using Grubbs' catalyst (8.1 mg, 0.1 equivalents, 0.05 equivalents per olefin) to yield polymer **12** in 50% yield:  $^1\text{H}$  NMR ( $\text{CDCl}_3$ )  $\delta$  7.6 (m), 7.3 (m), 6.2 (m), 6.1 (br s), 5.8 (br m), 3.4 (br m), 3.0 (br m), 2.8 (br m), 2.0 (br m). No GPC could be collected as the



polymer shows limited solubility in appropriate solvents. Polymer 12 exhibited a color change from colorless to purple upon irradiation with 310-nm light.

#### Example 4 – Preparation of Phenyl Homopolymer 17 (Figure 5)

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##### **Step 1 – Synthesis of phenyl derivative 14**

A mixture of toluene (10 mL) and aqueous Na<sub>2</sub>CO<sub>3</sub> (2 M, 5 mL) was degassed using argon for 30 minutes after which the mono-iodophotochrome 13 (583 mg, 1.39 mmol) and Pd(Ph<sub>3</sub>)<sub>4</sub> (5 mg) was added. Phenyl boronic acid (186 mg, 1.53 mmol) in dry ethanol (2 mL) was added and the mixture was heated at reflux for 24 hours. The reaction mixture was cooled to room temperature, extracted with ether (3 x 20 mL) and dried over Na<sub>2</sub>SO<sub>4</sub>. Purification via column chromatography through silica (hexane – 1:1 hexane-chloroform gradient) afforded 471 mg (92%) pure product as a pale pink solid: <sup>1</sup>H NMR (CDCl<sub>3</sub>) δ 7.47 (m, 2H), 7.32 (m, 2H), 7.21 (m, 1H), 6.97 (s, 1H), 6.61 (s, 1H), 2.75 (m, 4H), 2.05 (m, 5H), 1.87 (s, 3H); EIMS: 370.1 (M, 100%).

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##### **Step 2 – Synthesis of phenyl carboxylic acid 15**

A solution of chloro-phenylphotochrome 14 (450 mg, 1.2 mmol) in dry THF (20 mL) at -78°C under argon was treated with *tert*-butyllithium (0.7 mL of 1.7 M solution in hexane, 1.2 mmol). After stirring for 15 minutes, excess dry CO<sub>2</sub> was bubbled through the solution. The reaction mixture was warmed to room temperature, quenched with dilute HCl, extracted with CH<sub>2</sub>Cl<sub>2</sub> (3 x 50 mL), dried over Na<sub>2</sub>SO<sub>4</sub> and evaporated *in vacuo*. Purification by column chromatography through silica (3% CH<sub>3</sub>OH-CH<sub>2</sub>Cl<sub>2</sub>) afforded 365 mg (80%) of the product as a pale pink solid: <sup>1</sup>H NMR (CDCl<sub>3</sub>) δ 7.64 (s, 1H), 7.48 (m, 2H), 7.29 (m, 2H), 7.21 (m, 1H), 6.95 (s, 1H), 2.80 (m, 4H), 2.04 (m, 2H), 1.97 (s, 3H), 1.94 (s, 3H); <sup>13</sup>C NMR (CDCl<sub>3</sub>) δ 167.27, 144.93, 140.19, 137.50, 136.24, 136.20, 136.11, 134.48, 134.37, 133.43, 128.85, 128.16, 127.14, 125.39, 123.72, 38.63, 38.61, 27.04, 22.96, 15.02, 14.46; EIMS: 380.1 (M, 100%); IR (KBr-cast): 2952 (broad, m), 1669 (s).

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**Step 3 – Synthesis of phenyl-monomer 16**

A vigorously stirred solution of carboxylic acid 15 (301 mg, 0.79 mmol) and 6 drops of DMF in CH<sub>2</sub>Cl<sub>2</sub> (10 mL) at 0°C was treated with a solution of oxalyl chloride (502 mg, 3.95 mmol) in CH<sub>2</sub>Cl<sub>2</sub> (5 mL) dropwise over 10 minutes. After stirring at room temperature under argon for 2 hours, the reaction mixture was concentrated to dryness *in vacuo*. The residue was dissolved in CH<sub>2</sub>Cl<sub>2</sub> (10 mL) and added dropwise over 10 minutes to a solution of olefin 2 (305 mg, 1.19 mmol) and triethylamine (0.5 mL) in acetone cooled to 0°C. The mixture was stirred overnight under argon and the solvent was removed *in vacuo*. Purification by column chromatography through silica (2% CH<sub>3</sub>OH-CHCl<sub>3</sub>) afforded 472 mg (96%) of the product as a pale pink solid: <sup>1</sup>H NMR (CDCl<sub>3</sub>) δ 7.73 (s, 1H), 7.47 (m, 2H), 7.31 (m, 2H), 7.22 (m, 1H), 6.97 (s, 1H), 6.55 (m, 2H), 5.38 (m, 2H), 2.98 (m, 2H), 2.84 (m, 4H), 2.10 (m, 2H), 1.99 (s, 3H), 1.98 (s, 3H).

**Step 4 – Polymerization of phenyl-bicyclic olefin 16 to yield polymer 17**

The procedure is identical to that of monomer 5. Thus 16 (110 mg, 0.18 mmol) was polymerized with Grubbs' catalyst (5.85 mg, 0.04 equivalents) affording polymer 17 in 68% yield. <sup>1</sup>H NMR (CDCl<sub>3</sub>) δ 7.7 (br s), 7.4 (br s), 7.2 (br s), 6.95 (br s), 6.1 (br s), 5.8 (br d), 5.2 (br s), 4.6 (br d), 3.4 (br s), 2.8 (br s), 1.9 (m). GPC: Mn=11846, Mw=17536, Polydispersity Index=1.48. Polymer 17 exhibited a color change from colorless to purple upon irradiation with 254-nm light.

The polymers described in this application lend support to the mildness of the ROMP process. The selective ring-opening of the strained olefin in the monomer should be emphasized. As will be apparent from the above examples, the ROMP process produces homopolymers rather than copolymers. This provides macroscopic materials that have an increased density of photochromic units, a feature that is very advantageous for practical applications. Another advantage of using the ROMP technology is that by varying the catalyst-to-substrate stoichiometry, ROMP allows the polymer

chain length to be readily tailored. For example, extremely dense glass-like polymers having a very large number of photochromic units ( $n \geq 100$ ) could be prepared.

5           While the present invention has been described with reference to what are presently considered to be preferred examples, it is to be understood that the invention is not limited to the disclosed examples. On the contrary, the invention is intended to cover various modifications and  
10           equivalent arrangements included within the spirit and scope of the appended claims.

          All publications, patents and patent applications are herein incorporated by reference in their entirety to the same extent as if each individual publication, patent or patent application was specifically and  
15           individually indicated to be incorporated by reference in its entirety.

**FULL CITATIONS FOR REFERENCES REFERRED TO IN THE SPECIFICATION**

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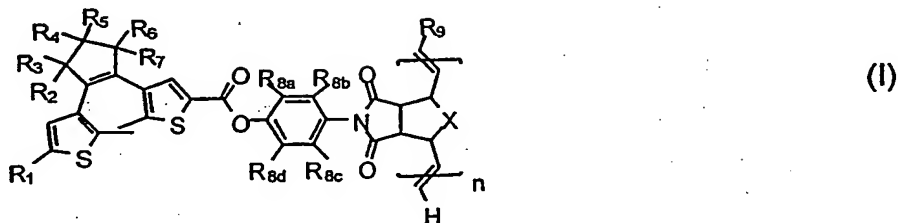
**TABLE 1****POLYMER YIELDS AND CHARACTERIZATION**

Photochrome	Equivalents monomer	Yield	M <sub>n</sub>	M <sub>w</sub>	M <sub>w</sub> /M <sub>n</sub>	λ <sub>max</sub> (nm) open form	λ <sub>max</sub> (nm) closed form
1a	25	76	7920	11917	1.50	248	512
1b	50	78	16771	24577	1.47	248	512
1c	100	75	28007	43820	1.56	248	512
5	-	-	-	-	-	248	512
9	25	53	7437	10953	1.47	250	559
8	-	-	-	-	-	250	559
12	10	50	-	-	-	266	562
11	-	-	-	-	-	266	562
17	25	68	11846	17536	1.48	270	553
16	-	-	-	-	-	270	553

SUBSTITUTE SHEET (RULE 26)

**What is claimed is:**

1. A homopolymer of the Formula I:

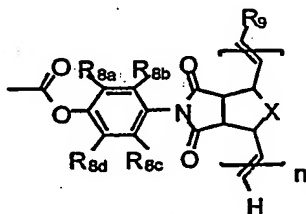


wherein:

$n$  is any integer;

$X$  is  $\text{CH}_2$ ,  $\text{CHR}_{10}$ ,  $\text{O}$ ,  $\text{S}$ ,  $\text{NH}$ ,  $\text{N-alkyl}$ ,  $\text{N-aryl}$ , or  $\text{CH}_2\text{-CH}_2$ ;

$R_1$  is selected from the group consisting of  $\text{H}$ , a halogen, a carboxylic acid, an ester, an aldehyde, an alkene, a phenyl, an aryl and structure A, where structure A is:



$R_2$ ,  $R_3$ ,  $R_4$ ,  $R_5$ ,  $R_6$  and  $R_7$  are each selected from the group consisting of  $\text{H}$  and a halogen;

$R_{8a}$ ,  $R_{8b}$ ,  $R_{8c}$  and  $R_{8d}$  are each selected from the group consisting of  $\text{H}$ , an alkyl and a halogen;

$R_9$  is a phenyl or an aryl; and,

R<sub>10</sub> is an alkyl, an aryl or a phenyl.

2. A homopolymer according to claim 1, wherein R<sub>1</sub> is chlorine, phenyl or structure A.

3. A homopolymer according to claim 1, wherein R<sub>2</sub>, R<sub>3</sub>, R<sub>4</sub>, R<sub>5</sub>, R<sub>6</sub> and R<sub>7</sub> are H or fluorine.

4. A homopolymer according to claim 1, wherein X is CH<sub>2</sub> or O.

5. A homopolymer according to claim 1, wherein R<sub>1</sub> is structure A and the homopolymer is cross-linked.

6. A homopolymer according to claim 1 or 2, wherein n is from about 10 to about 100.

7. A homopolymer according to claim 1 or 2, wherein n is 25.

8. A homopolymer according to claim 1 or 2, wherein n is 50

9. A homopolymer according to claim 1 or 2, wherein n is 100.

10. A method of preparing a homopolymer according to 1 comprising carrying out the reaction steps set forth in any one of Figures 1, 3, 4 and 5.

11. A method of preparing a photochromic homopolymer using ring-opening metathesis polymerization of a photochromic monomer.

12. The use of Grubbs' catalyst to catalyze the polymerization of a photochromic monomer to yield a photochromic homopolymer.



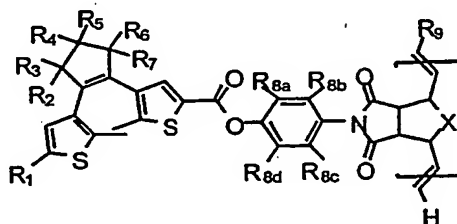
13. The use of claim 12, wherein said photochromic monomer is 1,2-bis-(3-thienyl)-cyclopentene.

14. A use of a homopolymer according to any one of claims 1 to 9.

15. A use according to claim 14 which is selected from the group consisting of:

- (1) ophthalmic lenses-eyeglasses that change color depending on the ambient light;
- (2) actinometry, and molecular sensors;
- (3) novelty items such as photochromic inks, paints and fibers;
- (4) variable transmission filters - those that on command, regulate the amount and type of light that can be transmitted;
- (5) high-density optical information storage systems (this invention is particularly well-suited to this application as it provides more information storage sites per unit area),
- (6) photo-regulated molecular switches that can be incorporated into molecular-scale machinery;
- (7) optoelectronic systems; and
- (8) reversible holographic systems.

16. A polymer comprising a compound of Formula VI:

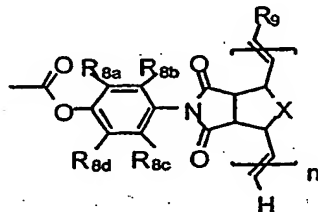


(VI)

wherein:

X is CH<sub>2</sub>, CHR<sub>10</sub>, O, S, NH, N-alkyl, N-aryl, or CH<sub>2</sub>-CH<sub>2</sub>;

$R_1$  is selected from the group consisting of H, a halogen, a carboxylic acid, an ester, an aldehyde, an alkene, a phenyl, an aryl and structure A, where structure A is:



$R_2$ ,  $R_3$ ,  $R_4$ ,  $R_5$ ,  $R_6$  and  $R_7$  are each selected from the group consisting of H and a halogen;

$R_{8a}$ ,  $R_{8b}$ ,  $R_{8c}$ , and  $R_{8d}$  are each selected from the group consisting of H, an alkyl and a halogen;

$R_9$  is a phenyl or an aryl; and,

$R_{10}$  is an alkyl, an aryl or a phenyl.

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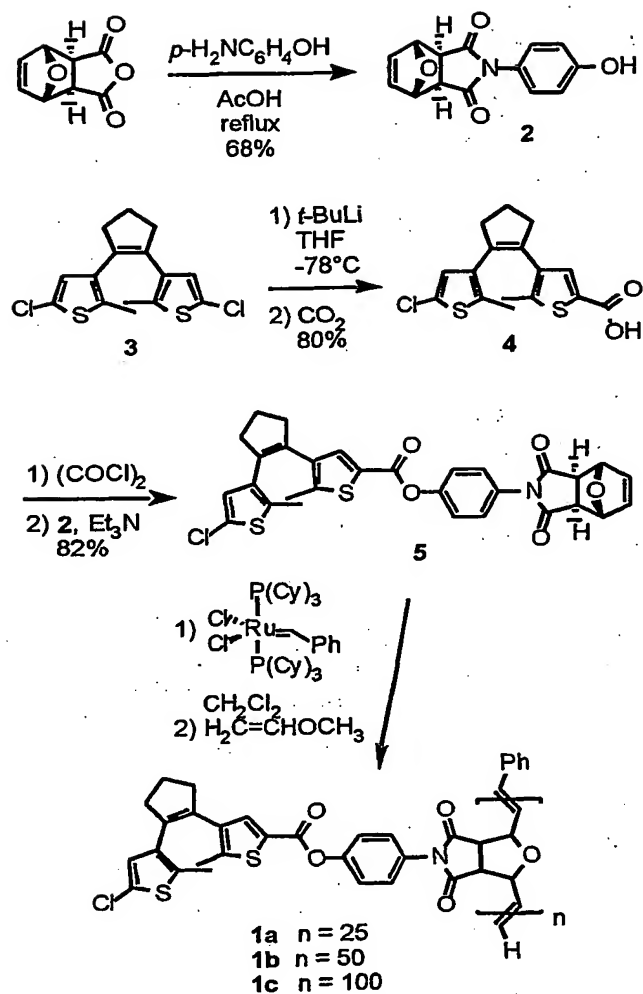


Figure 1 – Preparation of Homopolymers 1a-1c

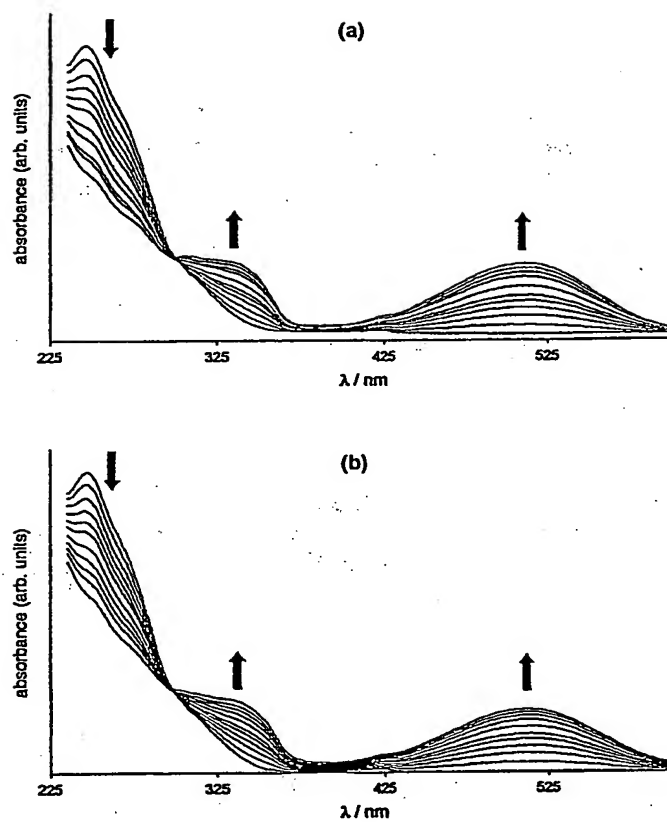


Figure 2

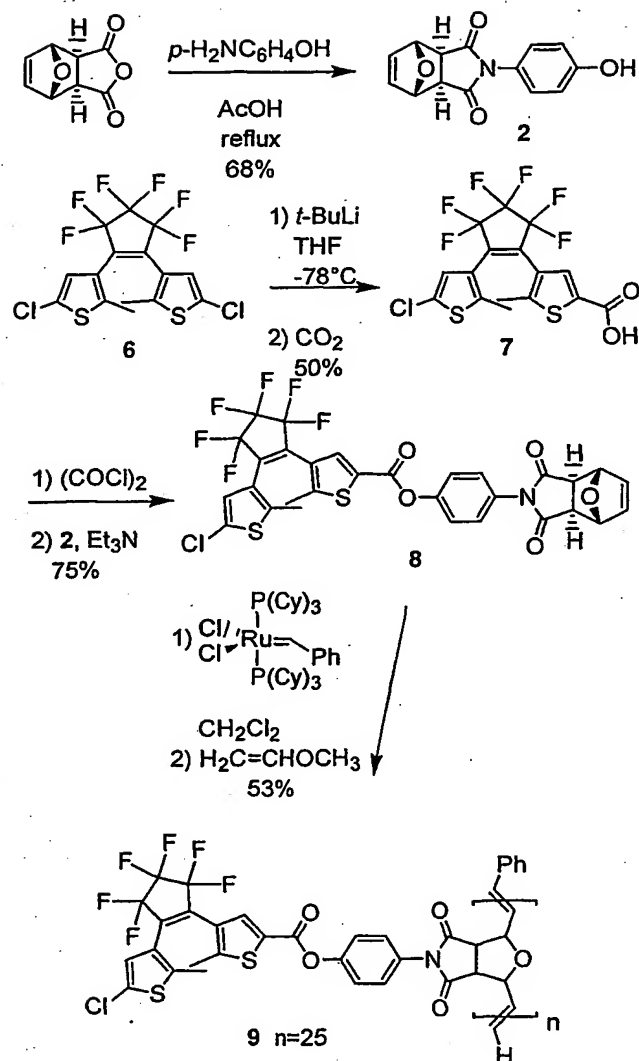


Figure 3 – Preparation of Fluorinated Homopolymer 9

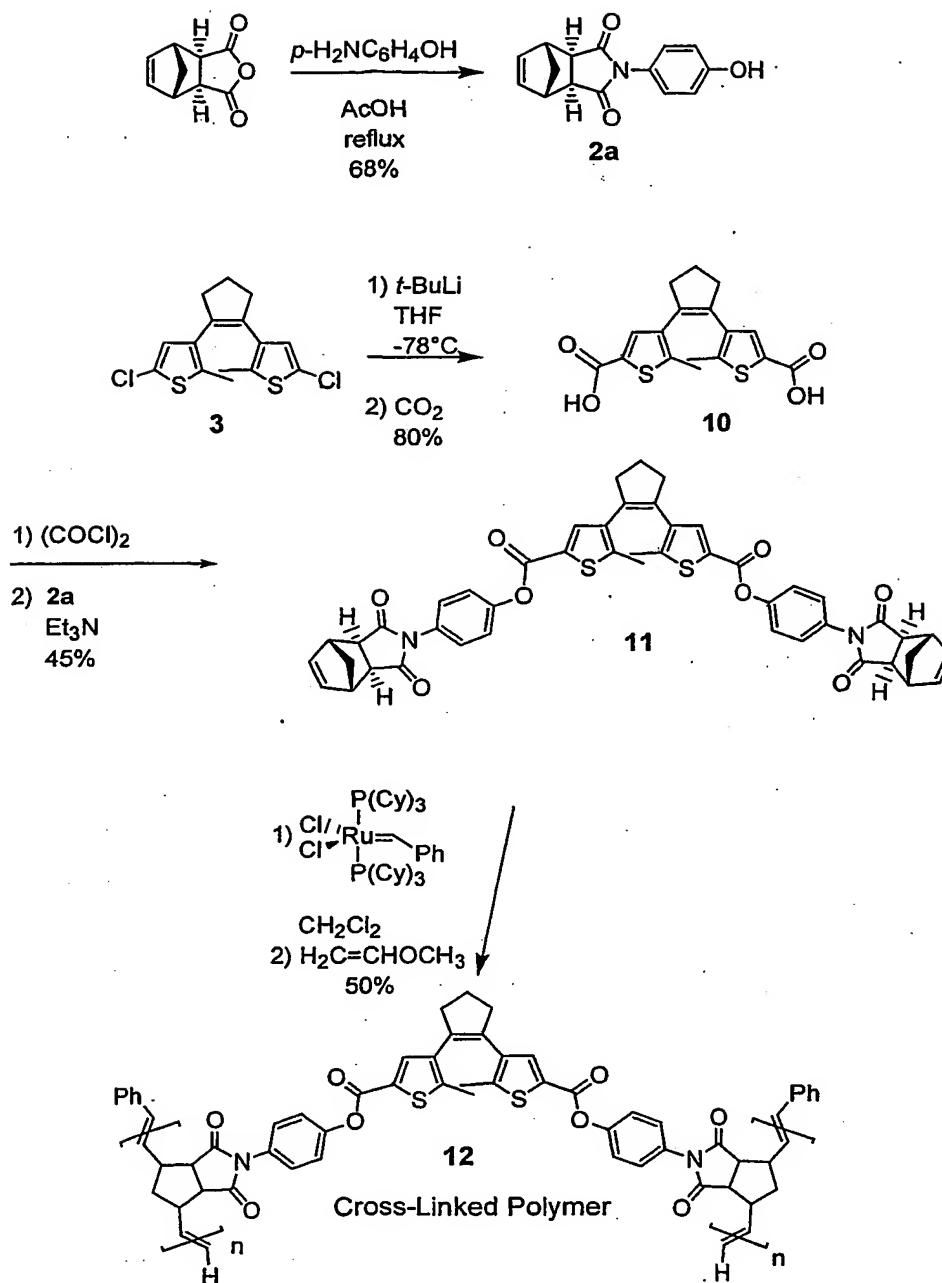


Figure 4 – Preparation of Cross-linked Homopolymer 12

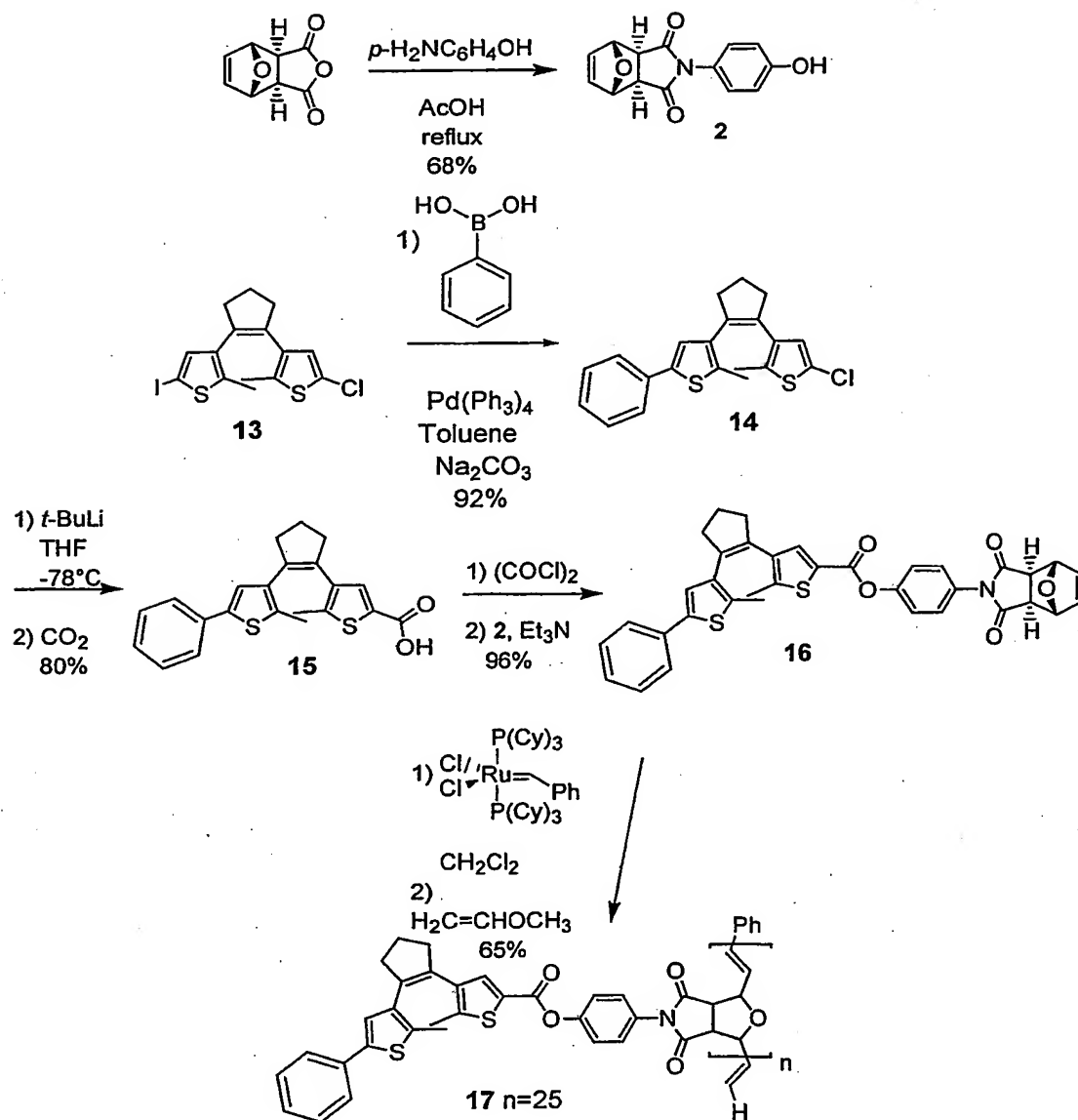


Figure 5 – Preparation of Phenyl Homopolymer 17

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(54) Title: NOVEL PHOTOCHROMIC POLYMERS AND METHODS OF SYNTHESIZING SAME

(57) Abstract: Ring-opening metathesis polymerization (ROMP) of a photochromic 1,2-bis-(3-thienyl)-cyclopentene monomer generated a series of novel polymers. All polymers exhibit reversible light-activated interconversion between their colorless-open and their colored-closed forms.

WO 02/06361 A3



<b>A. CLASSIFICATION OF SUBJECT MATTER</b> IPC 7 C08G61/12 C08G61/02		
According to International Patent Classification (IPC) or to both national classification and IPC		
<b>B. FIELDS SEARCHED</b> Minimum documentation searched (classification system followed by classification symbols) IPC 7 C08G C08F C07D		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
Electronic data base consulted during the international search (name of data base and, where practical, search terms used) EPO-Internal		
<b>C. DOCUMENTS CONSIDERED TO BE RELEVANT</b>		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5 342 909 A (GRUBBS ROBERT H ET AL) 30 August 1994 (1994-08-30) claim 1	12
A	KIM E ET AL: "PHOTOINDUCED REFRACTIVE INDEX CHANGE OF A PHOTOCHROMIC DIARYLETHENE POLYMER" MACROMOLECULES, AMERICAN CHEMICAL SOCIETY. EASTON, US, vol. 32, no. 15, 27 July 1999 (1999-07-27), pages 4855-4860, XP000845935 ISSN: 0024-9297 page 4857; figure 2 --- -/--	1-16
<input checked="" type="checkbox"/> Further documents are listed in the continuation of box C. <input checked="" type="checkbox"/> Patent family members are listed in annex.		
<b>* Special categories of cited documents:</b> *A* document defining the general state of the art which is not considered to be of particular relevance *E* earlier document but published on or after the international filing date *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) *O* document referring to an oral disclosure, use, exhibition or other means *P* document published prior to the international filing date but later than the priority date claimed *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art. *&* document member of the same patent family		
Date of the actual completion of the international search 11 January 2002		Date of mailing of the international search report 24/01/2002
Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016		Authorized officer Paalman, R

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